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Substitute for form 1449A/PTO				<i>Complete if Known</i>	
<b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>				Application Number	10/573,890
				Filing Date	03-29-06
				First Named Inventor	Kazutaka Nakamoto
				Art Unit	1625
				Examiner Name	Patricia L. Morris
				Attorney Docket Number	3939-0118PUS1
Sheet	1	of	3		

## U.S. PATENT DOCUMENTS

## FOREIGN PATENT DOCUMENTS

Examiner Initial *	Cite No. 1	Foreign Patent Document		Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, columns, Lines, Where Relevant Passages or Relevant Figures Appear	T 6
		Country <sup>3</sup> Code	Number <sup>4</sup> Kind Code (if known) <sup>5</sup>				
12	EP	0 124 154		11-07-1984			
	WO	03-091226		11-06-2003			
	WO	03-091227		11-06-2003			
	JP	2001-527083		12-25-2001			X
	WO	2004-014366		02-19-2004			
	JP	2005-526751		09-08-2005			X
	JP	2006-519247		08-24-2006			X
	WO	2009-084621		07-09-2009			X
	JP	52-94935		11-09-1993			X
	JP	59-073575		04-25-1984			X

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3. Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3) 4. For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. 5. Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.f 16 if possible. 6. Applicant is to place a check mark here if English language Translation is attached.

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<p>Substitute for form 1449A/PTO</p> <p><b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b></p> <p><i>(Use as many sheets as necessary)</i></p>				<p><b>Complete if Known</b></p>	
Sheet	2	of	3	Application Number	10/573,890
				Filing Date	03-29-06
				First Named Inventor	Kazutaka Nakamoto
				Art Unit	1625
				Examiner Name	Patricia L. Morris
				Attorney Docket Number	3939-0118PUS1

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3. Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3) 4. For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. 5. Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.16 if possible. 6. Applicant is to place a check mark here if English language Translation is attached.

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<b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>				Application Number	10/573,890
				Filing Date	03-29-06
				First Named Inventor	Kazutaka Nakamoto
				Group Art Unit	
				Examiner Name	
Sheet	3	of	3	Attorney Docket Number	3939-0118PUS1

<b>NON PATENT LITERATURE DOCUMENTS</b>				
Examiner initial *	Cite No. 1	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.		T <sup>2</sup>
	23	CHANG et al., "Synthesis and Structure-Activity Relationships of Quaternary Ammonium Cephalosporins with 3-Pyrazolylpyridinium Derivatives," <i>Bioorganic &amp; Medicinal Chemistry Letters</i> (2000) Vol. 10, No. 11, pp. 1211-1214		
	24	CONNORS et al., "Prodrugs in medicine," <i>Overview, Biologicals &amp; Immunologicals, Exp. Opin. Ther. Patents</i> , Vol. 5, No. 9, 1995, pp. 873-885		
	25	Copy of an Office Action from co-pending U.S. Patent Application No. 11/589,128, mailed May 7, 2009		
	26	HATA, "New Approaches to Antifungal Drugs for the Treatment of Fungal and Protozoal Infections, Ravaconazole and Beyond: New Targets and Pre-clinical Strategies," <i>The SMI's 12th Annual Conference, Superbugs and Superdrugs</i> , March 18, 2010, Crowne Plaza London - St. James, 44 pages		
	27	International Search Report dated May 20, 2008 for corresponding International Application No. PCT/JP2008/057851		
	28	ISHIKAWA et al., "TAK-599, a Novel N-Phosphono Type Prodrug of Anti-MRSA Cephalosporin T-91825: Synthesis, Physicochemical and Pharmacological Properties," <i>Bioorganic &amp; Medicinal Chemistry</i> , Vol. 11, pp. 2427-2437, (2003)		
	29	LUKEVICS et al., "Synthesis and cytotoxicity of silyl- and carbonyl-substituted isoxazoles," <i>Chemistry of Heterocyclic Compounds</i> (2000) Vol. 36, No. 10, pp. 1226-1231		
	30	PLATE et al., "Synthesis and Muscarinic Activities of 3-(Pyrazolyl)-1,2,5,6-tetrahydropyridine Derivatives," <i>Bioorganic &amp; Medicinal Chemistry Letters</i> (1996) Vol. 4, No. 2, pp. 227-237		
	31	Supplementary European Search Report dated February 6, 2009 for corresponding European Application No. 04788159.4		
	32	TANAKA et al., "An Effective Lewis Acid-Mediated 1,3-Dipolar Cycloaddition of Nitrile Oxide Using Acetylene: Synthesis of a (2-Aminopyridin-3-yl) isoxazole Derivative and Its Application to Novel Antifungal Agents," pp. 1-8		
	33	VRZHESCHCH et al., "Supercooperativity in platelet aggregation: Substituted pyridyl isoxazoles, a new class of supercooperative platelet aggregation inhibitors," <i>FEBS Letters</i> (1994) Vol. 351, No. 2, pp. 168-170		

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